AMENDMENTS TO THE SPECIFICATION:

Please delete the paragraph beginning at page 23, line 25 and replace it with the following paragraph:

-- and wherein any CH_2 or CH_3 group within a R^1 substituent optionally bears on each said CH_2 or CH_3 group a substituent selected from hydroxy, amino, (1-6C)alkoxy, (1-6C)alkylsulphonyl, (1-6C)alkylamino and di-[(1-6C)alkyl]amino, or from a group of the formula:

$$-X^3-Q^5$$

wherein X^3 is a direct bond or is selected from O, **N(R^7)** ~~**N(R^6)**~~, $CON(R^7)$, $N(R^7)CO$ and $C(R^7)_2O$, wherein R^7 is hydrogen or (1-6C)alkyl, and Q^5 is heteroaryl, heteroaryl-(1-6C)alkyl, heterocyclyl or heterocyclyl-(1-6C)alkyl, --

Please delete the paragraph beginning at page 24, line 31 and replace it with the following paragraph:

-- and wherein any CH_2 or CH_3 group within a R^1 substituent optionally bears on each said CH_2 or CH_3 group a substituent selected from hydroxy, amino, (1-6C)alkoxy, (1-6C)alkylsulphonyl, (1-6C)alkylamino and di-[(1-6C)alkyl]amino, or from a group of the formula:

$$-X^3-Q^5$$

wherein X^3 is a direct bond or is selected from O, **N(R^7)** ~~**N(R^6)**~~, $CON(R^7)$, $N(R^7)CO$ and $C(R^7)_2O$, wherein R^7 is hydrogen or (1-6C)alkyl, and Q^5 is heteroaryl, heteroaryl-(1-6C)alkyl, heterocyclyl or heterocyclyl-(1-6C)alkyl, --

Please delete the paragraph beginning at page 76, line 16 and replace it with the following paragraph:

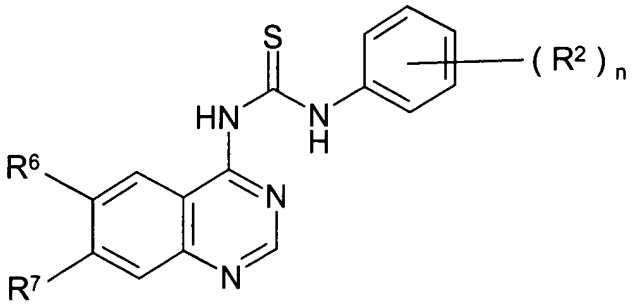
-- A mixture of 7-benzyloxy-6-methoxy-3,4-dihydroquinazolin-4-one (25.1 g), thionyl chloride (450 ml) and DMF (1 ml) was stirred and heated to reflux for 2 hours. The mixture was evaporated and the residue was dissolved in toluene and the solution was evaporated. The resultant solid was suspended in methylene chloride (500 ml) , solid potassium carbonate (39 g) was added and the mixture was stirred for 10 minutes. Water (500 ml) was added and the

mixture stirred for another 10 minutes. The methylene chloride layer was separated, dried over magnesium sulphate and evaporated. The residue was purified by column chromatography on silica using increasingly polar mixtures of methylene chloride and ethyl acetate as eluent. There was thus obtained **7-benzyloxy-4-chloro-6-methoxyquinazoline** ~~**7-benzyl-4-chloro-6-methoxyquinazoline**~~ (21.54 g); NMR Spectrum: ($DMSOd_6$) 4.0 (s, 3H), 5.36 (s, 2H), 7.31-7.46 (m, 4H), 7.51 (d, 2H), 7.58 (s, 1H), 8.88 (s, 1H). --

Please delete the paragraph beginning at page 137, line 6 and replace it with the following paragraph:

-- Using an analogous procedure to that described in Example 27, the appropriate 4-aminoquinazoline was reacted with the appropriate isothiocyanate to give the compounds described in Table VII.

Table VII



No.	R^6	R^7	$(R^2)_n$	Note
1	methoxy	N-methylpiperidin-4-ylmethoxy	2,6-dichloro	(a)
2	methoxy	N-methylpiperidin-4-ylmethoxy	2,6-difluoro	(b)
3	methoxy	N-methylpiperidin-4-ylmethoxy	2-chloro-6-methyl	(c)
4	methoxy	N-methylpiperidin-4-ylmethoxy	2,4,6-trichloro	(d)
5	methoxy	N-methylpiperidin-4-ylmethoxy	2,6-dimethyl-4-bromo	(e)
6	methoxy	N-methylpiperidin-4-ylmethoxy	2,5-dimethyl	(f)
7	methoxy	3-pyrrolidin-1-ylpropoxy	2,6-dichloro	(g)
8	methoxy	3-pyrrolidin-1-ylpropoxy	2,6-difluoro	(h)
9	methoxy	3-pyrrolidin-1-ylpropoxy	2-chloro-6-methyl	(i)

10	methoxy	2-(2-methoxyethoxy)ethoxy	2,6-dimethyl	(j)
11	methoxy	2-morpholinoethoxy	2,6-dimethyl	(k)
12	methoxy	3-morpholinopropoxy	2,6-dimethyl	(l)
13	methoxy	cyclopropylmethoxy	2,6-dimethyl	(m)
14	methoxy	2-morpholinoethoxy	**2-chloro-6-methyl** ~~**2-chloro-6-dimethyl**~~	(n)
15	methoxy	3-morpholinopropoxy	2-chloro-6-methyl	(o)
16	methoxy	N-methylpiperidin-4-ylmethoxy	2-methyl	(p)
17	methoxy	2-pyrrolidin-1-ylethoxy	2,6-dimethyl	(q)

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